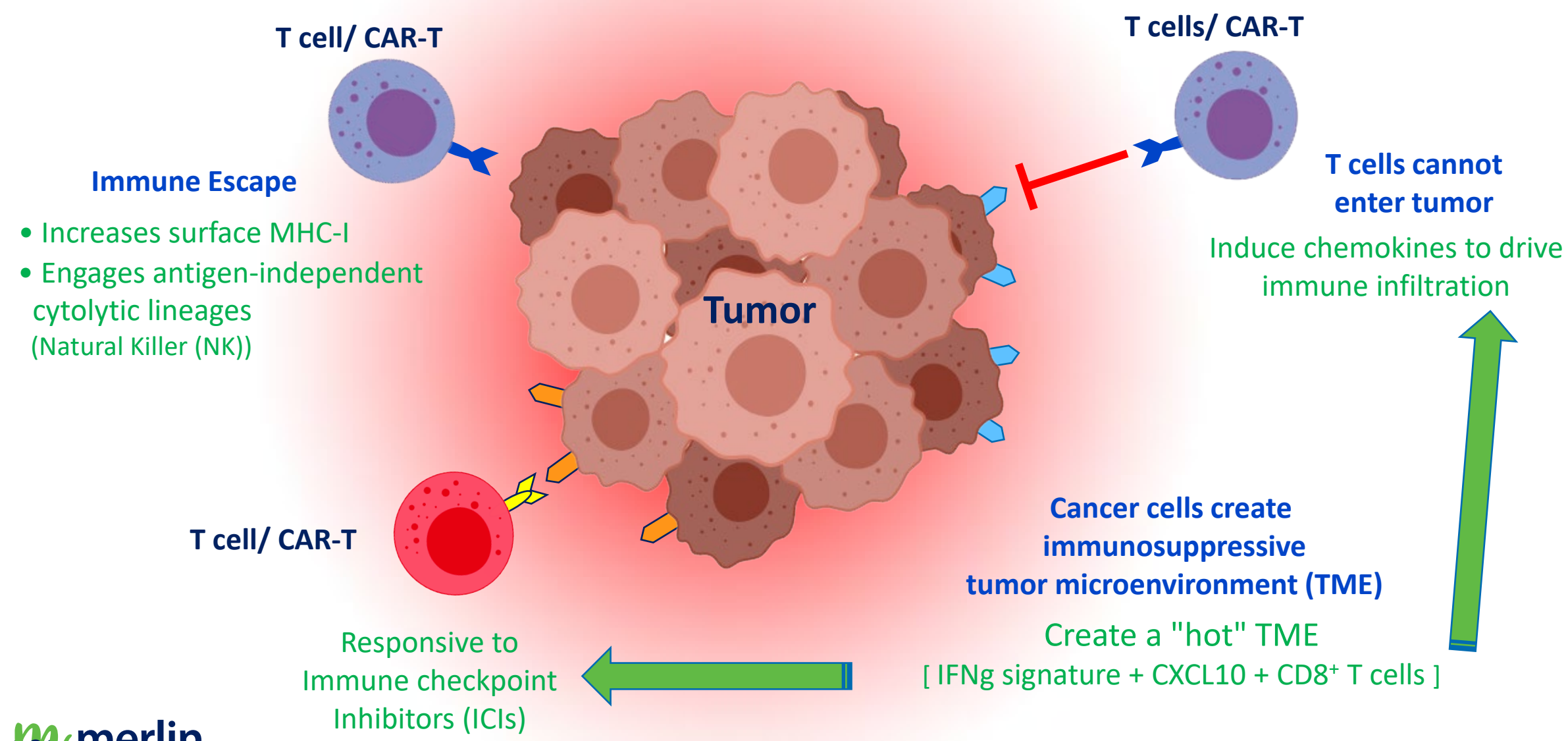


**RANDALL N.
HYER
CEO**

June 2023



MER-101 (USP6) ACTIVATES SEVERAL IMMUNO-STIMULATORY PATHWAYS



EPIDEMIOLOGICAL DATA VALIDATES USP6 AS A UNIQUE IMMUNOTHERAPY TARGET

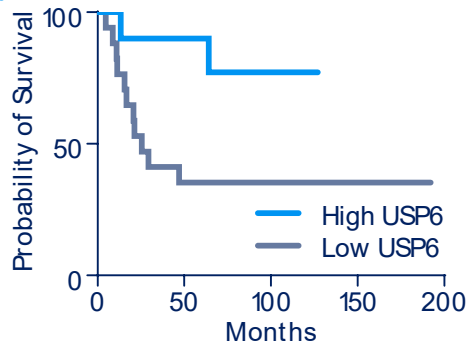


USP6

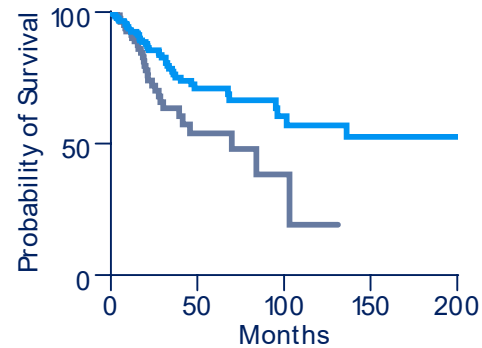
Real world expression data



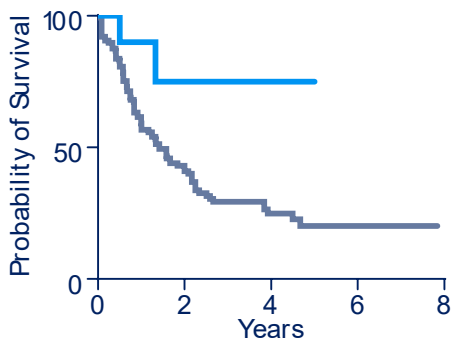
Ewing sarcoma:



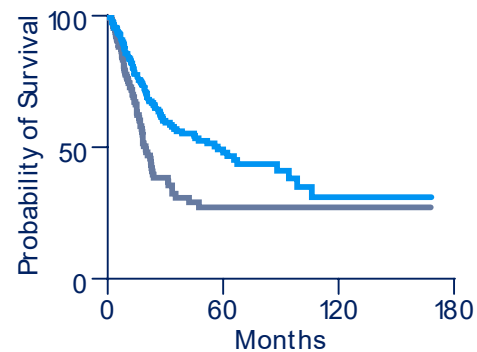
Cervical cancer:



Acute myeloid leukemia (AML):



Bladder cancer:



WHY USP6?



Elevated USP6 tumor expression is beneficial for patients

Correlates with enhanced immune activation and improved outcomes



Unique, multi-pronged mechanism of action

Activation of multiple immunostimulatory pathways reduces likelihood of resistance



Novel target, first-in-class therapy

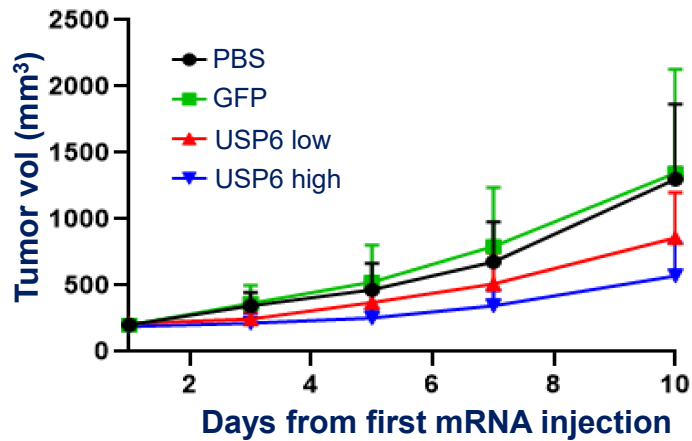
Only company seeking to exploit USP6's powerful anti-tumor activity

Proof of principle

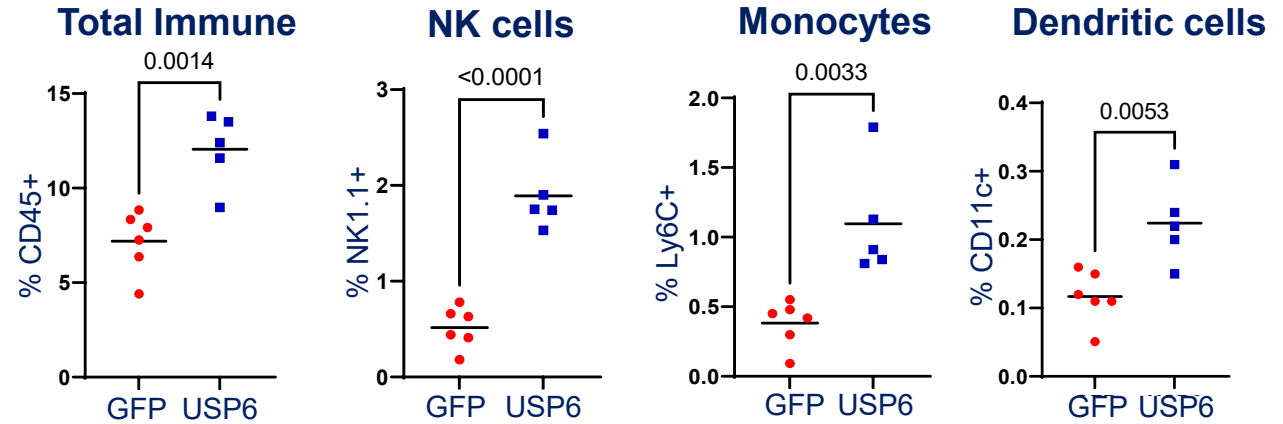
Intratumoral USP6 mRNA suppresses Ewing sarcoma tumorigenesis and triggers intratumoral and systemic immune activation



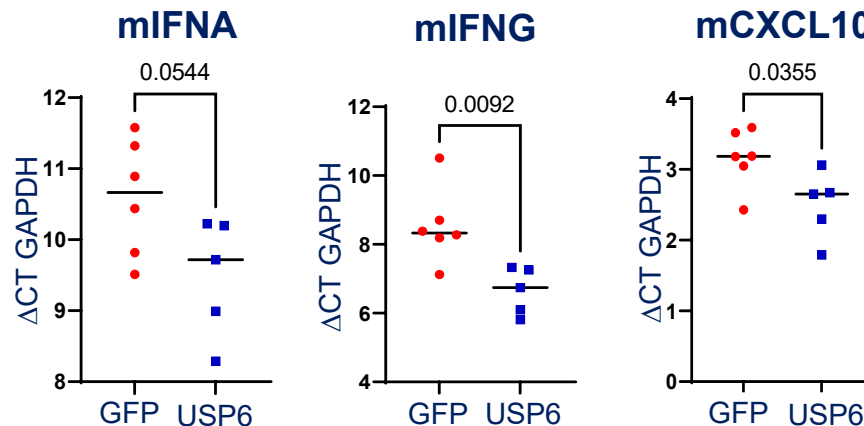
USP6 mRNA
Lipid nanoparticles (LNPs)
Intratumoral delivery



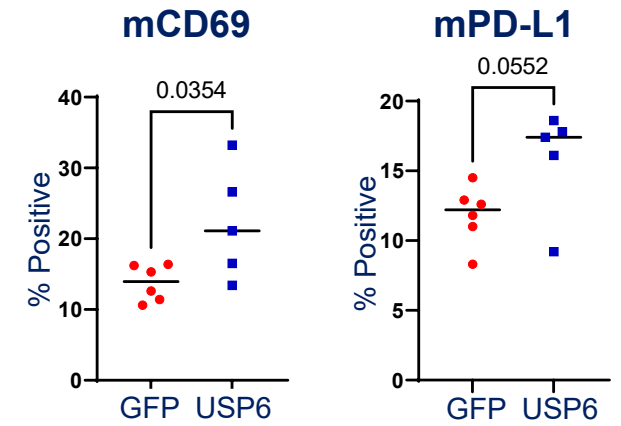
Intratumoral immune infiltration



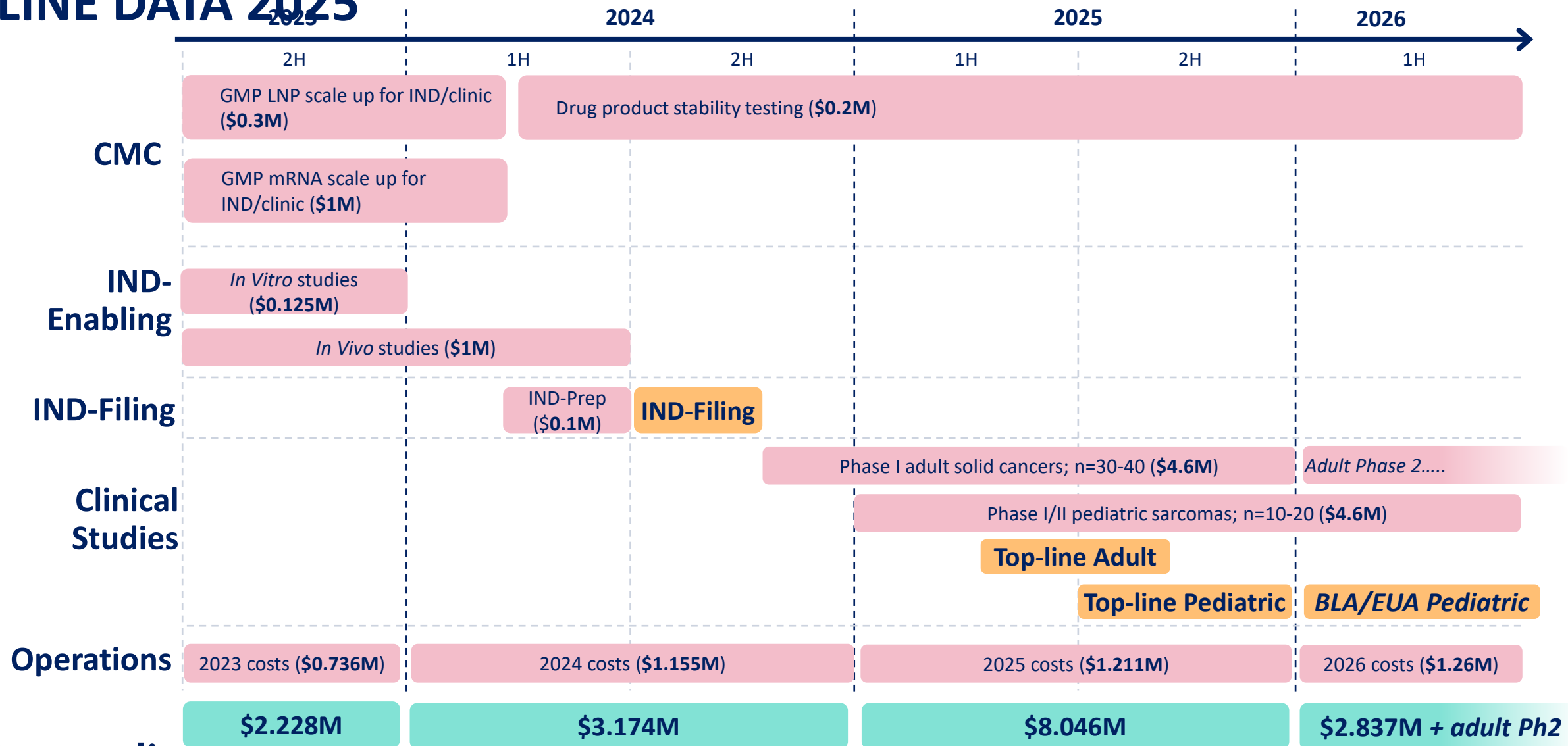
Intratumoral immune activation



Systemic (splenic) NK activation



FDA GUIDANCE ENABLES CLINICAL ENTRY IN 2024 WITH TOP LINE DATA 2025



MER-101 HAS LARGE MARKET SIZE DRIVEN BY POTENTIAL APPLICATION TO NUMEROUS CANCERS AND EXPANSION OPPORTUNITIES

US Revenue Model*

	2030	2034
Solid Tumor Market	130,000,000,000	199,000,000,000
Immunotherapy Capture	63,000,000,000	95,000,000,000
Est. VA Price/Cycle	9,400,000,000	14,300,000,000
Cycles/Yearly	15,500	18,100
Market Capture	668,000	869,000
Yearly Revenue	1%	5%
	\$103,000,000	\$787,000,000

5 year cumulative

\$2,069,287,658

Upside opportunities

- Market penetration
- COGS reduction
- Pricing
- Tier increase

MER-101 broad immuno-modulator:

- ✓ Numerous potential indications.
- ✓ Expansion to adjuvant/neo-adjuvant.
- ✓ Expansion to combination with other immunotherapies.

INVESTMENT SUMMARY

Lead Asset MER-101:

- ✓ **Novel, 1st-in-class** cancer immunotherapy with **exclusive IP in major world-wide markets.**
- ✓ **Demonstrated efficacy** in multiple mouse models with **supporting human epidemiological survival data.**
- ✓ Data supports **cancer agnostic indication** with an adult injectable tumor market conservative initial 5-year sales cumulative estimate at **~\$2 billion.**
- ✓ Development in pediatric sarcomas can **expedite approvals with a Ph1/2 readout in 2026 H1** with eligibility for a **~\$100M+ priority review voucher.**

Financials and timeline:

Current Funding



\$650K

Pennsylvania state grant
(**Non-dilutive**)



\$105K

SAFE Notes

Seeking



Tranche of \$20M

\$5M

to advance MER-101 to IND
(target 2024 H1)

\$15M

to generate MER-101 Phase 1/2 data
(target: 2026 H1)